091823733

FILE 'HOME' ENTERED AT 10:50:21 ON 29 APR 2002

=> file reg

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

0.21

0.21

FILE 'REGISTRY' ENTERED AT 10:50:32 ON 29 APR 2002 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2002 American Chemical Society (ACS)

28 APR 2002 HIGHEST RN 408492-65-9 STRUCTURE FILE UPDATES: DICTIONARY FILE UPDATES: 28 APR 2002 HIGHEST RN 408492-65-9

TSCA INFORMATION NOW CURRENT THROUGH July 7, 2001

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Calculated physical property data is now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details: http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf

*** YOU HAVE NEW MAIL ***

Uploading 09823283.str

L1 STRUCTURE UPLOADED

=> d l1

L1 HAS NO ANSWERS

L1 STR

Structure attributes must be viewed using STN Express query preparation.

=> s l1 full

FULL SEARCH INITIATED 10:50:56 FILE 'REGISTRY' 999 TO ITERATE FULL SCREEN SEARCH COMPLETED -

100.0% PROCESSED 999 ITERATIONS 9 ANSWERS

SEARCH TIME: 00.00.01

9 SEA SSS FUL L1 L_2

=> file caplus COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY SESSION 140.28 140.49 FULL ESTIMATED COST

FILE 'CAPLUS' ENTERED AT 10:51:09 ON 29 APR 2002 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2002 AMERICAN CHEMICAL SOCIETY (ACS)

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FILE COVERS 1907 - 29 Apr 2002 VOL 136 ISS 18 FILE LAST UPDATED: 28 Apr 2002 (20020428/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

CAS roles have been modified effective December 16, 2001. Please check your SDI profiles to see if they need to be revised. For information on CAS roles, enter HELP ROLES at an arrow prompt or use the CAS Roles thesaurus (/RL field) in this file.

*** YOU HAVE NEW MAIL ***

=> s 11

REG1stRY INITIATED

Substance data SEARCH and crossover from CAS REGISTRY in progress... Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

SAMPLE SEARCH INITIATED 10:51:21 FILE 'REGISTRY' SAMPLE SCREEN SEARCH COMPLETED - 43 TO ITERATE

43 ITERATIONS 100.0% PROCESSED

1 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE** BATCH **COMPLETE**

PROJECTED ITERATIONS: 467 TO 1253

PROJECTED ANSWERS: 1 TO

1 SEA SSS SAM L1 L3

1 L3 L4

=> s 12 7 L2 L5

=> d 17 bib abs hitstr 1-7

L7 NOT FOUND

The L-number entered has not been defined in this session, or it

has been deleted. To see the L-numbers currently defined in this session, enter DISPLAY HISTORY at an arrow prompt (=>).

=> d 15 bib abs hitstr 1-7

ANSWER 1 OF 7 CAPLUS COPYRIGHT 2002 ACS 1.5

2002:107826 CAPLUS AN

DN 136:172758

Terminally-branched polymeric linkers containing extension moieties for ΤI prodrug conjugates

IN Greenwald, Richard B.; Choe, Yun H.

PΑ USA

SO U.S. Pat. Appl. Publ., 32 pp. CODEN: USXXCO

DTPatent

English LA

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PΙ	US 2002015691	A1	20020207	US 2001-823296	20010329
PRAI	US 2000-193931P	P	20000331		

The present invention relates to polymer-based (e.g., PEG) conjugates having increased therapeutic payloads. In particular, the invention relates to the use of extension moieties which increase the efficiency of the loading of drugs onto the polymeric carriers. A variety of prodrugs were prepd. from ara-C and PEG derivs. by using spacer groups. The prodrug demonstrated better antitumor activity than ara-C alone. The prodrug produced complete tumor regression.

396134-08-0P 396134-17-1P IΤ

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(terminally-branched polymeric linkers contg. extension moieties for prodrug conjugates)

396134-08-0 CAPLUS RN

Poly(oxy-1,2-ethanediyl), .alpha.-[18-[(1-.beta.-D-arabinofuranosyl-1,2-CN dihydro-2-oxo-4-pyrimidinyl)amino]-4,4-bis[14-[(1-.beta.-Darabinofuranosyl-1,2-dihydro-2-oxo-4-pyrimidinyl)amino]-3,14-dioxo-7,10dioxa-4,13-diazatetradec-1-yl]-2,7,18-trioxo-11,14-dioxa-3,8,17triazaoctadec-1-yl]-.omega.-methoxy- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-C

PAGE 3-B

RN 396134-17-1 CAPLUS

CN Poly(oxy-1,2-ethanediyl), .alpha.-methyl-.omega.-hydroxy-, 1-ether with N-(hydroxyacetyl)-L-aspartoylbis[N1,N4-bis[2-[2-[2-[[[(1-.beta.-D-arabinofuranosyl-1,2-dihydro-2-oxo-4-pyrimidinyl)amino]carbonyl]amino]ethoxy]ethoxy]ethyl]-L-aspartamide] (9CI) (CA INDEX NAME)

PAGE 2-A

$$\begin{array}{c} \mathsf{CH}_2 \\ \mathsf{CH}_2 \\ \mathsf{C} \\ \mathsf{C} \\ \mathsf{CH}_2 \\ \mathsf{CH}_2 \\ \mathsf{CH}_2 \\ \mathsf{NH} \\ \mathsf{C} \\$$

PAGE 2-C

$$-\operatorname{CH}_2-\operatorname{O}-\operatorname{CH}_2-\operatorname{CH}_2-\operatorname{NH}-\operatorname{C}-\operatorname{NH}$$

PAGE 4-B

```
ANSWER 2 OF 7 CAPLUS COPYRIGHT 2002 ACS
L5
     2001:75282 CAPLUS
AN
DN
     134:136702
     Enhanced circulation effector composition and method
ΤI
     Zalipsky, Samuel; Woodle, Martin C.; Martin, Francis J.; Barenholz,
IN
     Yechezkel
PΑ
     Sequus Pharmaceuticals, Inc., USA
    U.S., 32 pp., Cont.-in-part of U.S. Ser. No. 316,436, abandoned.
SO
     CODEN: USXXAM
DT
    Patent
LA
    English
FAN.CNT 2
```

PATENT NO. KIND DATE

APPLICATION NO. DATE

ΡI		6180134 6326353	B1 B1	20010130 20011204		1995-480332 1993-35443	19950607 19930323
	US	2001043929	A1	20011122	US	2001-877978	20010608
PRAI	US	1993-35443	A2	19930323			
	US	1994-316436	B2	19940929			
	US	1995-480332	A 1	19950607			

AB A liposome compn. comprising small, surface-bound effector mols. is disclosed. The liposomes have a surface layer of hydrophilic polymer chains, for enhanced circulation time in the bloodstream. The effector mols. are attached to the distal ends of the polymer chains. In one embodiment, the effector is polymyxin B, for treatment of septic shock. Liposomes with covalently bound .beta.- galactosidase were prepd. from a maleimide deriv. of distearyl phosphatidyl ethanolamine carbamide of PEG bis(amine), .alpha.-tocopherol, cholesterol, partially hydrogenated egg phosphatidylcholine, egg phosphatidyl glycerol, and .beta.-galactosidase.

IT 159125-99-2P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(prepn. of polyethylene glycol derivs. for liposomes contg. polypeptides or polysaccharide effector mols. covalently attached therewith)

RN 159125-99-2 CAPLUS

CN Poly(oxy-1,2-ethanediyl), .alpha.-[2-[[(2Z)-3-carboxy-1-oxo-2-propenyl]amino]ethyl]-.omega.-[[9-hydroxy-9-oxido-4,15-dioxo-12-[(1-oxooctadecyl)oxy]-8,10,14-trioxa-3,5-diaza-9-phosphadotriacont-1-yl]oxy]-(9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

IT 159158-15-3P

RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of polyethylene glycol derivs. for liposomes contg. polypeptides or polysaccharide effector mols. covalently attached therewith)

RN 159158-15-3 CAPLUS

CN Poly(oxy-1,2-ethanediyl), .alpha.-[9-hydroxy-9-oxido-4,15-dioxo-12-[(1-oxooctadecyl)oxy]-8,10,14-trioxa-3,5-diaza-9-phosphadotriacont-1-yl].omega.-[2-[[1-oxo-3-(2-pyridinyldithio)propyl]amino]ethoxy]- (9CI) (CA INDEX NAME)

PAGE 1-C

- (CH₂)₁₆-Me

RE.CNT 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 3 OF 7 CAPLUS COPYRIGHT 2002 ACS

AN 1999:78360 CAPLUS

DN 130:175340

TI Thermal printing material containing phenolic compound as color developer

IN Shimada, Masaru; Matsui, Hiroaki; Torii, Masaaki

PA Ricoh Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 39 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN CNT 1

PAN.	CNTT				
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PΙ	JP 11028868	A2	19990202	JP 1997-197918	19970708
OS	MARPAT 130:17534	0			
GI					

- * STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY AVAILABLE VIA OFFLINE PRINT *
- AB The title material comprises a support coated with a heat-sensitive layer contg. a leuco dye and, as a color developer, a phenolic compd. selected from I-VI (X = hydrocarbon, alkoxy, halo, H; n = 1-3; m = 0-2; p = 0-2 in I and II, p = 1-10 in III-VI; q1, q2, q3 = 1-10; q4 = 0-21). The material provides high d. and low fog images with good plasticizer resistance and water resistance.
- IT 220427-37-2 220427-38-3 220427-40-7

220427-42-9

RL: TEM (Technical or engineered material use); USES (Uses) (thermal printing material contg. phenolic compd. as color developer)

RN 220427-37-2 CAPLUS

CN 5,8-Dioxa-2,11-diazatridecanamide, 13-(4-hydroxyphenyl)-N-octadecyl-12-oxo-(9CI) (CA INDEX NAME)

PAGE 1-B

- (CH₂)₁₇ - Me

RN 220427-38-3 CAPLUS

CN 5,8-Dioxa-2,11-diazatetradecanamide, 14-(4-hydroxyphenyl)-N-octadecyl-12-oxo-(9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 220427-40-7 CAPLUS

CN 5,8-Dioxa-2,11-diazatridecanamide, 13-[(4-hydroxyphenyl)amino]-N-octadecyl-12-oxo-(9CI) (CA INDEX NAME)

- NH- (CH₂)₁₇- Me

220427-42-9 CAPLUS RN

5,8-Dioxa-2,11-diazatetradec-13-enamide, 14-(4-hydroxyphenyl)-N-octadecyl-CN12-oxo- (9CI) (CA INDEX NAME)

PAGE 1-A

$$\begin{array}{c} \text{CH} & \text{CH} & \text{CH} - \text{CH} - \text{CH}_2 - \text{CH}_2$$

PAGE 1-B

ANSWER 4 OF 7 CAPLUS COPYRIGHT 2002 ACS L5

1997:308339 CAPLUS ΑN

DN126:334416

Liposomes for treatment of B-cell and T-cell disorders ΤI

Allen, Theresa M.; Martin, Francis J. IN

Sequus Pharmaceuticals, Inc., USA PA

U.S., 31 pp. Cont.-in-part of U.S. 5,527,528. SO CODEN: USXXAM

DTPatent

English LA

FAN.C					
]	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI U	US 5620689	A	19970415	US 1995-475050	19950607
Ţ	US 5013556	Α	19910507	US 1989-425224	19891020
i	AU 9066374	A1	19910516	AU 1990-66374	19901019
i	AU 642679	B2	19931028		
]	EP 496813	A1	19920805	EP 1990-916409	19901019
]	EP 496813		19941214		
	R: AT, BE,	CH, DE	, DK, ES, FR,	GB, GR, IT, LI, LU	, NL, SE
	JP 05505173	T2	19930805	JP 1990-515238	19901019
	JP 2001181214	A2	20010703	JP 2001-4291	
1	US 5213804	A	19930525	US 1991-642321	
]	NO 9201213	Α	19920604		
	FI 9201763	A	19920421	FI 1992-1763	
1	US 5527528	Α	19960618	US 1993-40544	
	JP 10001431	A2	19980106	JP 1997-63661	19970317
	JP 2889549		19990510		
PRAI	US 1989-425224	A2	19891010		
1	US 1991-642321	A2	19910115		
1	US 1993-40544	A2	19930331		
1	JP 1990-515238	A3	19901019		

JP 1991-501034 A3 19901019 WO 1990-US6034 A 19901019

AB A method of treating a subject having a disorder characterized by a neoplasm of B-lymphocyte or T-lymphocyte lineage cells is described. The method includes administering a suspension of liposomes having a surface coating of polyethylene glycol chains. Attached to the distal ends of the chains are antibodies or antibody fragments effective to bind to an antigen specific to the affected cells. In one embodiment, anti-CD19 antibodies are attached to the liposome-bound chains, for treatment of multiple myeloma. An example PEG compd., distearoylphosphatidylethanolami ne PEG hydrazide deriv. was prepd.

IT 159158-15-3DP, reaction products with antibodies

159158-15-3P 179267-38-0P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation) (liposomes for treatment of B-cell and T-cell disorders)

RN 159158-15-3 CAPLUS

CN Poly(oxy-1,2-ethanediyl), .alpha.-[9-hydroxy-9-oxido-4,15-dioxo-12-[(1-oxooctadecyl)oxy]-8,10,14-trioxa-3,5-diaza-9-phosphadotriacont-1-yl].omega.-[2-[[1-oxo-3-(2-pyridinyldithio)propyl]amino]ethoxy]- (9CI) (CA INDEX NAME)

PAGE 1-A

DAGE 1-R

PAGE 1-C

- (CH₂)₁₆-Me

RN 159158-15-3 CAPLUS

CN Poly(oxy-1,2-ethanediyl), .alpha.-[9-hydroxy-9-oxido-4,15-dioxo-12-[(1-oxooctadecyl)oxy]-8,10,14-trioxa-3,5-diaza-9-phosphadotriacont-1-yl]-.omega.-[2-[[1-oxo-3-(2-pyridinyldithio)propyl]amino]ethoxy]- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

$$\begin{array}{c} O \\ | \\ | \\ | \\ -O \\ \hline \end{array} \\ \begin{array}{c} O \\ | \\ | \\ CH_2 - CH_2 - NH - C \\ -NH - CH_2 - CH_2 - O \\ -P - O - CH_2 - CH - CH_2 - O \\ -P \\ | \\ O \end{array} \\ \begin{array}{c} O \\ | \\ CH_2 - CH_2 - O \\ -P \\ O - CH_2 - CH_2 - O \\ -P - O - CH_2 - CH_2 - O \\ -P - O - CH_2 - CH_2 - O \\ -P - O - CH_2 - CH_2 - O \\ -P - O - CH_2 - CH_2 - O - C \\ -P - O - CH_2 - CH_2 - CH_2 - O - C \\ -P - O - CH_2 - CH_2 - CH_2 - O - C \\ -P - O - CH_2 - CH_2 - CH_2 - O - C \\ -P - O - CH_2 - CH_2 - CH_2 - CH_2 - CH_2 - O - C \\ -P - O - CH_2 - C$$

PAGE 1-C

$$-$$
 (CH₂)₁₆-Me

RN 179267-38-0 CAPLUS

Poly(oxy-1,2-ethanediyl), .alpha.-[2-[(3-carboxy-1-oxo-2-propenyl)amino]ethyl]-.omega.-[[9-hydroxy-9-oxido-4,15-dioxo-12-[(1-oxooctadecyl)oxy]-8,10,14-trioxa-3,5-diaza-9-phosphadotriacont-1-yl]oxy]-(9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

```
1996:449885 CAPLUS
ΑN
    125:105097
DN
    Solid tumor treatment method using antitumor agent-containing liposomes
ТT
    with PEG coating and surface-attached antibody
    Allen, Theresa M.; Martin, Francis J.
ΤN
    Sequus Pharmaceuticals, Inc., USA
PΑ
    U.S., 17 pp. Cont.-in-part of U.S. 5,213,804.
SO
    CODEN: USXXAM
DT
    Patent
    English
LA
FAN.CNT 9
     PATENT NO.
                   KIND DATE
                                       APPLICATION NO. DATE
                                         ______
                          _ _ _ _ _ _ _ _
     A
                                        US 1993-40544
                                                         19930331
                          19960618
PΙ
    US 5527528
                     Α
                          19910507
                                       US 1989-425224
                                                         19891020
    US 5013556
                     A1
                                        AU 1990-66374
    AU 9066374
                          19910516
                                                         19901019
                    В2
    AU 642679
                          19931028
                    A1
                                        EP 1990-916409
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    EP 496813
                          19920805
                    В1
    EP 496813
                         19941214
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE
     JP 05505173 T2 19930805 JP 1990-515238 19901019
                    A2
                                                         19901019
     JP 2001181214
                          20010703
                                        JP 2001-4291
    US 5213804
NO 9201213
                                       US 1991-642321
                                                        19910115
                          19930525
                     Α
                                       NO 1992-1213
                                                        19920327
                          19920604
                     A
     FI 9201763
                         19920421
                                        FI 1992-1763
                                                        19920421
                     Α
                                        WO 1994-US3457 19940330
     WO 9422429
                     A1 19941013
        W: AU, CA, JP, US
        RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE
                    Al 19941024 AU 1994-65272 19940330
     AU 9465272
     US 5620689
                     Α
                          19970415
                                        US 1995-475050
                                                        19950607
    JP 10001431
JP 2889549
                                        JP 1997-63661 19970317
                    A2 19980106
                    B2 19990510
PRAI US 1989-425224 A2 19891020
     US 1991-642321 A2 19910115
     JP 1990-515238 A3 19901019
     JP 1991-501034 A3 19901019
                          19901019
     WO 1990-US6034 A
                   A2
W
                         19930331
     US 1993-40544
     WO 1994-US3457
                          19940330
     A method of administering an antitumor compd. to a subject is disclosed.
AB
     The method includes administering liposomes having sizes predominantly in
     the range 0.05 to 0.12 .mu., and contg. an antitumor compd. in
     liposome-entrapped form, a surface coating of polyethylene glycol chains,
     at a surface concn. thereof sufficient to extend the blood circulation
     time of the liposomes severalfold over that of liposomes in the absence of
     such coating, and surface-attached antibody mols. effective to bind
     specifically to tumor-assocd. antigens present at the tumor site. One
     liposome compn. includes doxorubicin in entrapped form, and, on the
     liposome surface, a monoclonal antibody against highly proliferating cells
     in a lung squamous cell carcinoma.
     179267-38-0P
ΙT
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
        (prepn. and reaction; antitumor agent-contg. liposome prepn. with PEG
        coating and surface-attached antibody for solid tumor treatment)
     179267-38-0 CAPLUS
RN
     Poly(oxy-1,2-ethanediyl), .alpha.-[2-[(3-carboxy-1-oxo-2-
CN
     propenyl) amino] ethyl] - .omega. - [[9-hydroxy-9-oxido-4,15-dioxo-12-[(1-
     oxooctadecyl)oxy]-8,10,14-trioxa-3,5-diaza-9-phosphadotriacont-1-yl]oxy]-
```

(9CI) (CA INDEX NAME)

L5 ANSWER 6 OF 7 CAPLUS COPYRIGHT 2002 ACS

AN 1995:196581 CAPLUS

DN 122:38832

TI Pharmaceutical liposomes comprising PEG for administration of polypeptides

IN Zalipsky, Samuel; Martin, Francis

PA Liposome Technology, Inc., USA

SO PCT Int. Appl., 53 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

PΙ

PRAI

•	-14 T	_				
	PAT	TENT NO.	KIND	DATE	APPLICATION NO.	DATE
	WO	9421281	A1	19940929	WO 1994-US3102	19940322
		W: AU, CA,	JP			
		RW: AT, BE,	CH, DE	, DK, ES, FR,	GB, GR, IE, IT, LU	, MC, NL, PT, SE
	AU	9463683	A1	19941011	AU 1994-63683	19940322
[US	1993-35640		19930323		
	WO	1994-US3102		19940322		

AB Pharmaceutical liposomes comprising PEG are prepd. for administration of polypeptides. Liposomes contg. biotin-PEG were incubated in the presence of avidin. Avidin-coated liposomes were incubated with biotinylated IgG to obtain liposome-bound antibody.

IT 159125-99-2P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation)
 (pharmaceutical liposomes comprising PEG for administration of
 polypeptides)

RN 159125-99-2 CAPLUS

CN Poly(oxy-1,2-ethanediyl), .alpha.-[2-[[(2Z)-3-carboxy-1-oxo-2-propenyl]amino]ethyl]-.omega.-[[9-hydroxy-9-oxido-4,15-dioxo-12-[(1-oxooctadecyl)oxy]-8,10,14-trioxa-3,5-diaza-9-phosphadotriacont-1-yl]oxy]-(9CI) (CA INDEX NAME)

```
ANSWER 7 OF 7 CAPLUS COPYRIGHT 2002 ACS
L5
```

AN1994:686620 CAPLUS

DN 121:286620

Pharmaceutical liposomes comprising hydrophilic polymer conjugates with ΤI polypeptides or polysaccharides

IN Zalipsky, Samuel; Woodle, Martin C.; Martin, Francis J.; Barenholz, Yechezkel

PΑ Liposome Technology, Inc., USA

PCT Int. Appl., 72 pp. CODEN: PIXXD2

DT Patent

English

FAN.	CNT	2				
	PAT	TENT NO.	KIND	DATE	APPLICATION NO. DATE	
ΡI	WO	9421235		19940929	WO 1994-US3103 19940322	
		W: AU, CA, RW: AT, BE,		, DK, ES,	FR, GB, GR, IE, IT, LU, MC, NL, PT, SE	
	US	6326353	В1	20011204	US 1993-35443 19930323	
	CA	2157410	AA	19940929	CA 1994-2157410 19940322	
	ΑU	9463684	A1	19941011	AU 1994-63684 19940322	
	ΕP	689428	A1	19960103	EP 1994-910988 19940322	
	EΡ	689428	В1	19990120		
		R: AT, BE,	CH, DE	, DK, ES,	FR, GB, IE, IT, LI, LU, NL, SE	
	JР	08508256	T2	19960903	JP 1994-521332 19940322	
	AT	175868	E	19990215	AT 1994-910988 19940322	
	ES	2131190	Т3	19990716	ES 1994-910988 19940322	
PRAI	US	1993-35443	Α	19930323		
	WO	1994-US3103	W	19940322		

A liposome compn. comprising small, surface-bound effector mols., such as AΒ .beta.-galactosidase (I), is disclosed. The liposomes have a surface layer of hydrophilic polymer chains, such as PEG, for enhanced circulation time in the bloodstream. The effector mols. are attached to the distal ends of the polymer chains. Maleic acid deriv. of distearoylphosphatidylcholine (DSPE) carbamide of PEG bis(amine) was heated with acetic anhydride satd. with anhyd. NaCH3COO for 2 h at 50.degree. to obtain maleimide of DSPE carbamide of PEG bis(amine) (II) which was purified to a pale yellow, viscous oil. I liposomes with covalently-bound II was prepd.

IT 159125-99-2P 159158-15-3P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation) (pharmaceutical liposomes comprising hydrophilic polymer conjugates with polypeptides or polysaccharides)

RN 159125-99-2 CAPLUS

Poly(oxy-1,2-ethanediyl), .alpha.-[2-[[(2Z)-3-carboxy-1-oxo-2-propenyl]amino]ethyl]-.omega.-[[9-hydroxy-9-oxido-4,15-dioxo-12-[(1-oxooctadecyl)oxy]-8,10,14-trioxa-3,5-diaza-9-phosphadotriacont-1-yl]oxy]-(9CI) (CA INDEX NAME)

PAGE 1-A

$$_{
m HO_2C-CH-}$$
 $_{
m CH-}$ $_{
m CH-CH_2-CH_2-}$ $_{
m CH_2-CH_2-}$ $_{
m CH_2-CH_2-}$ $_{
m CH_2-CH_2-}$ $_{
m CH_2-CH_2-}$

PAGE 1-B

RN 159158-15-3 CAPLUS

CN Poly(oxy-1,2-ethanediyl), .alpha.-[9-hydroxy-9-oxido-4,15-dioxo-12-[(1-oxooctadecyl)oxy]-8,10,14-trioxa-3,5-diaza-9-phosphadotriacont-1-yl]-.omega.-[2-[[1-oxo-3-(2-pyridinyldithio)propyl]amino]ethoxy]- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

PAGE 1-C

- (CH₂)₁₆-Me

=>